

1/31

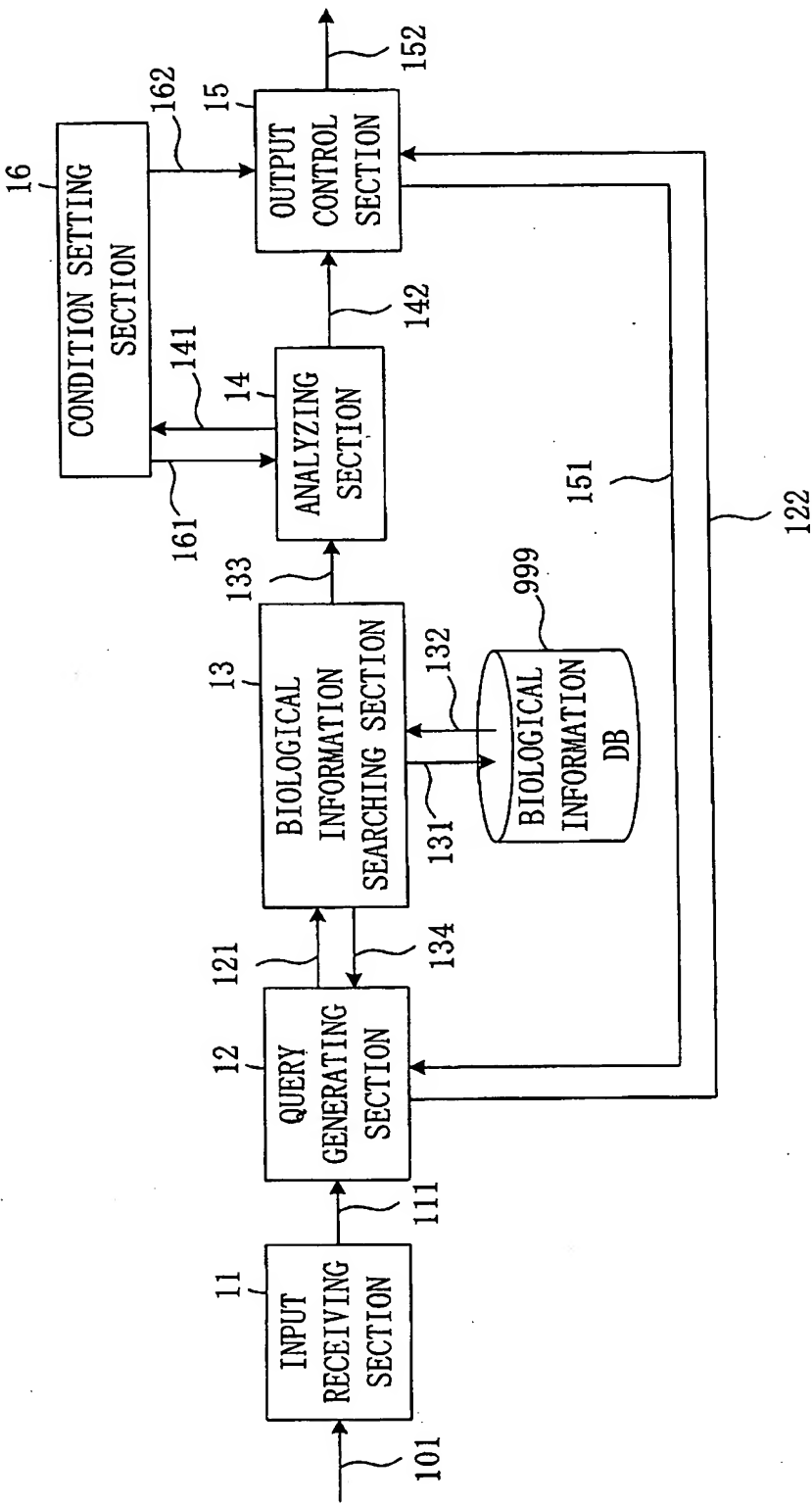


FIG. 1

2/31

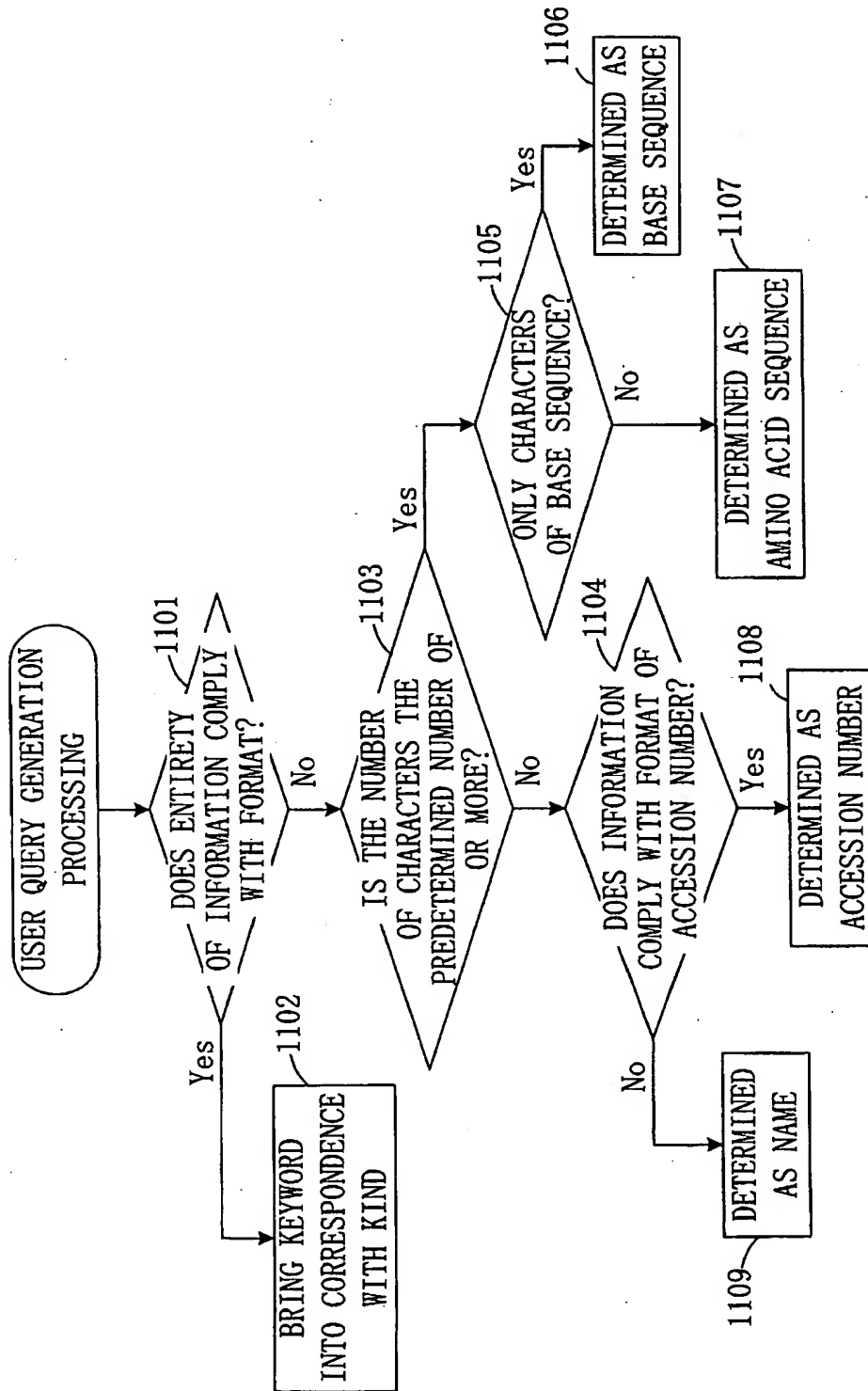


FIG. 2

3/31

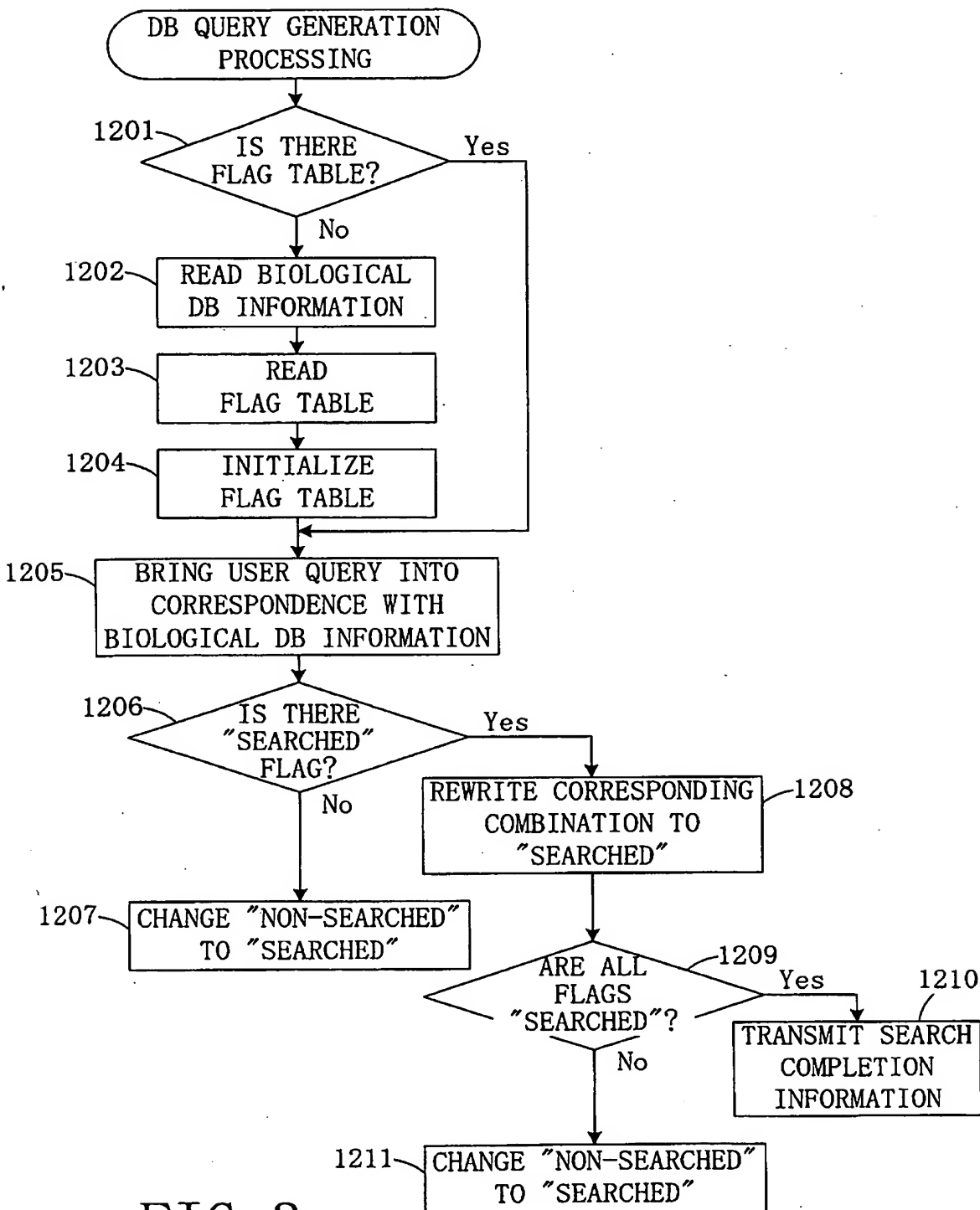


FIG. 3

4/31

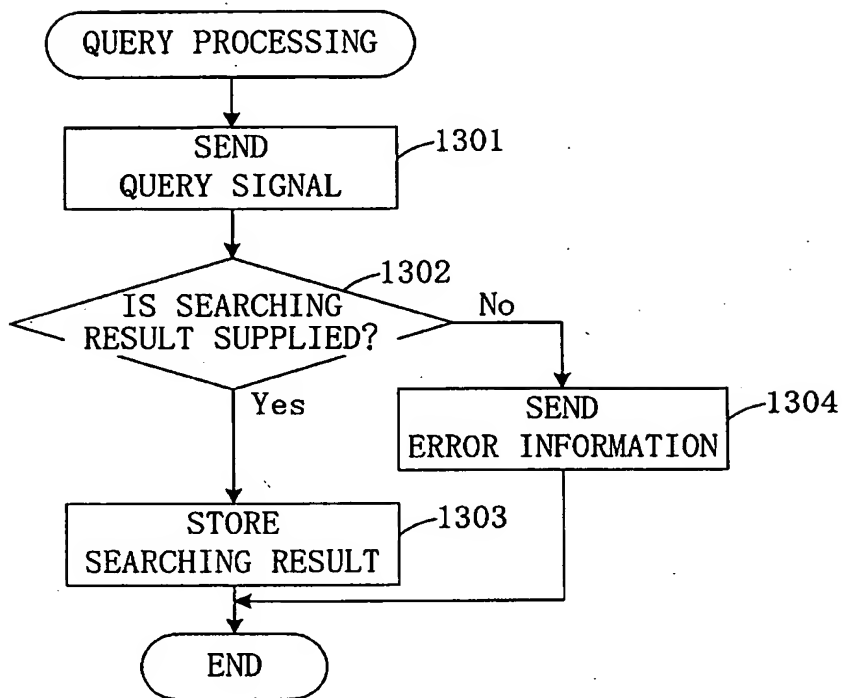


FIG. 4

5/31

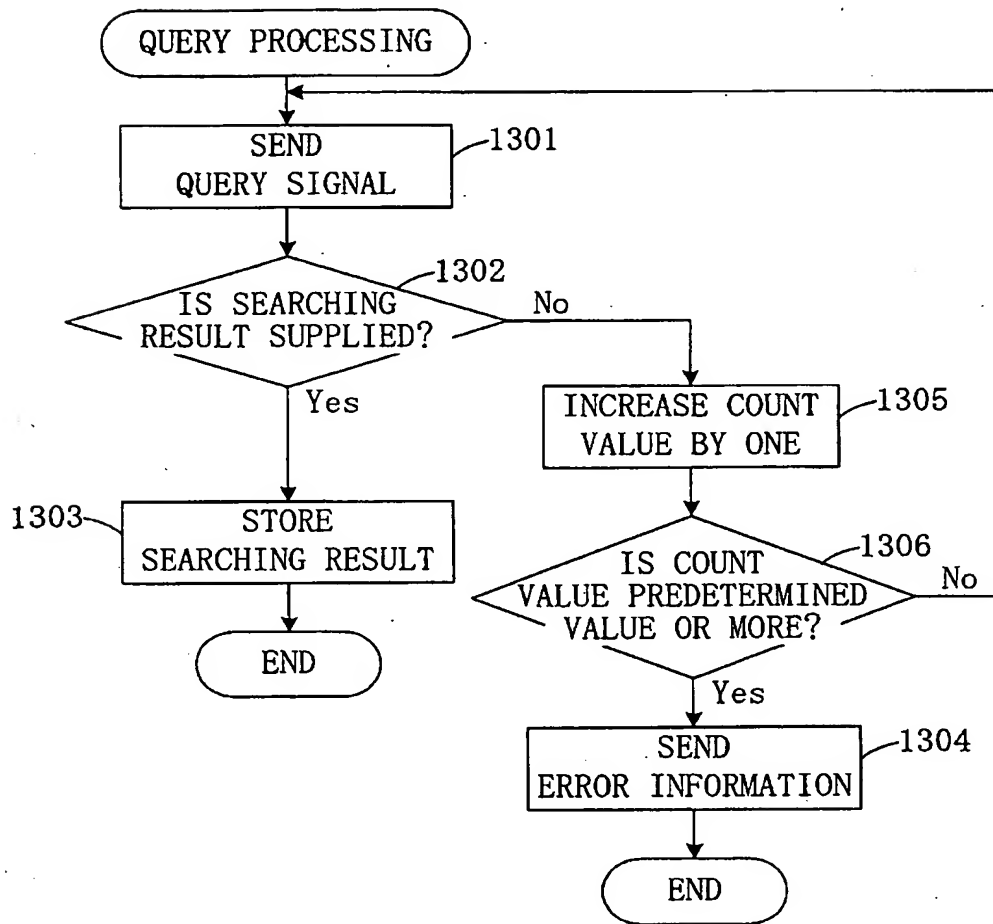


FIG. 5

6/31

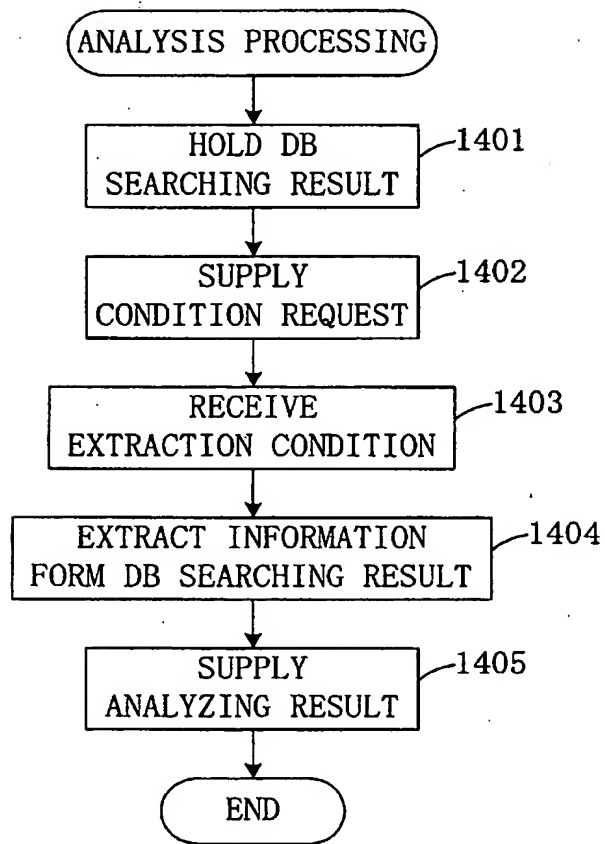


FIG. 6

7/31

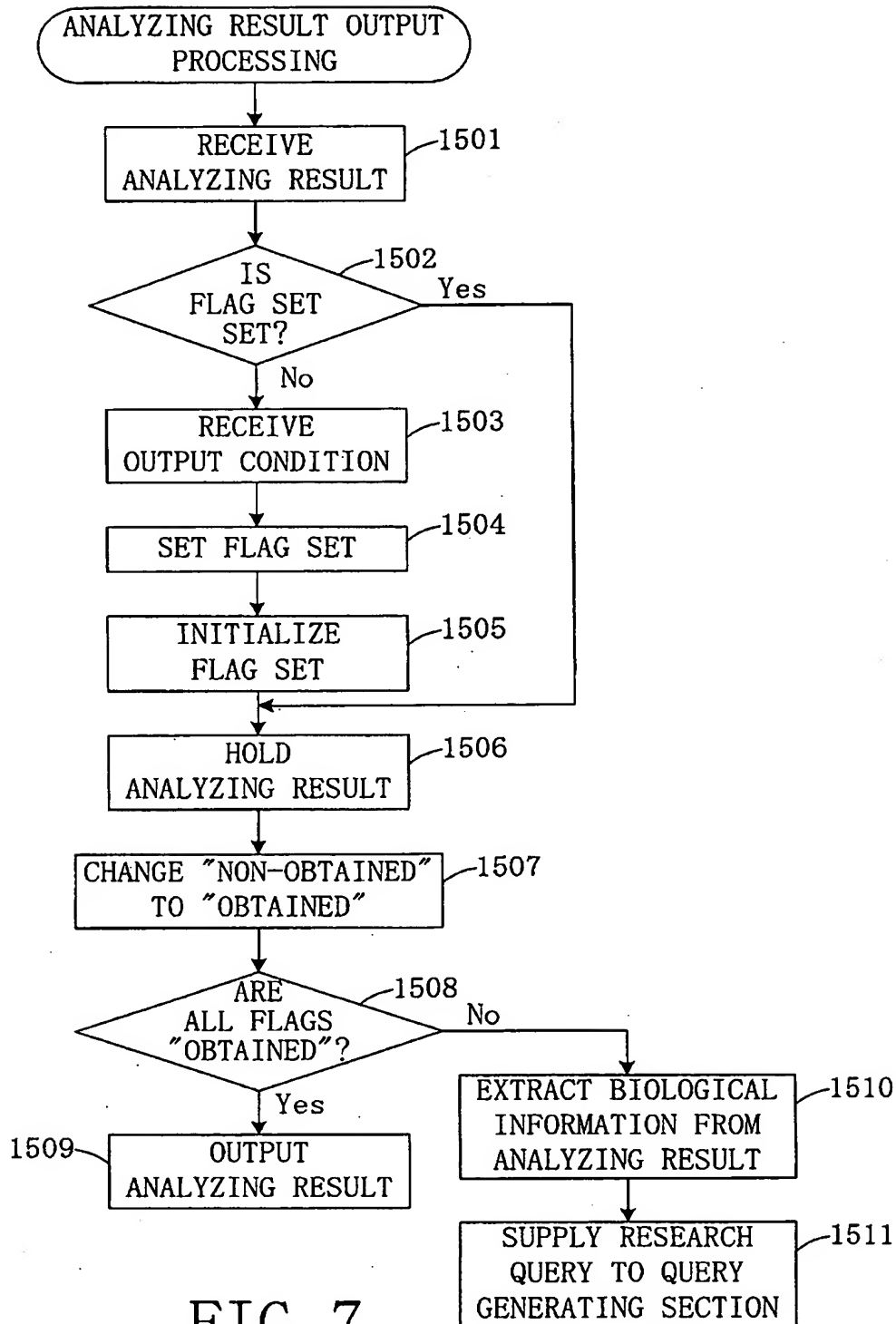


FIG. 7

8/31

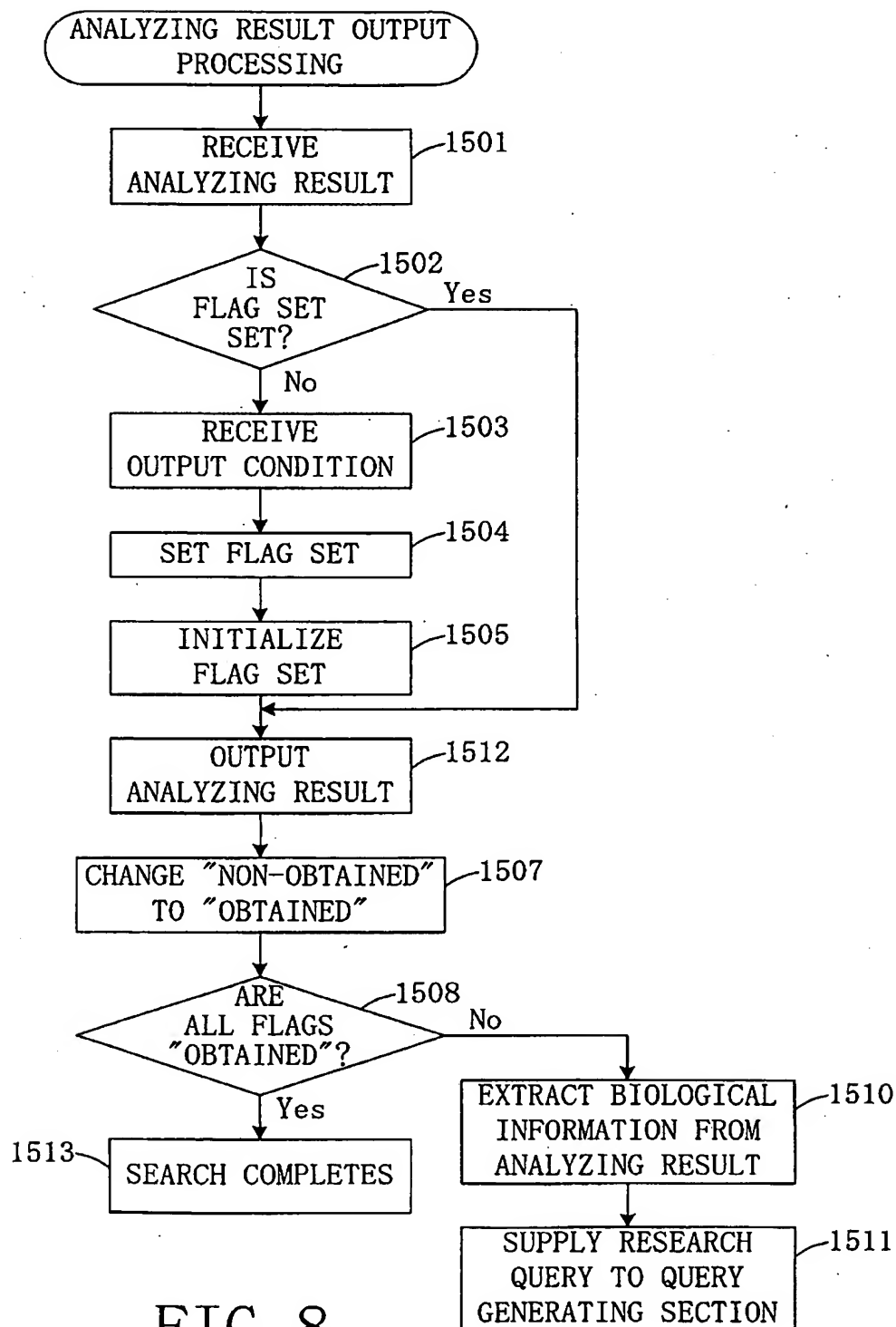


FIG. 8

9/31

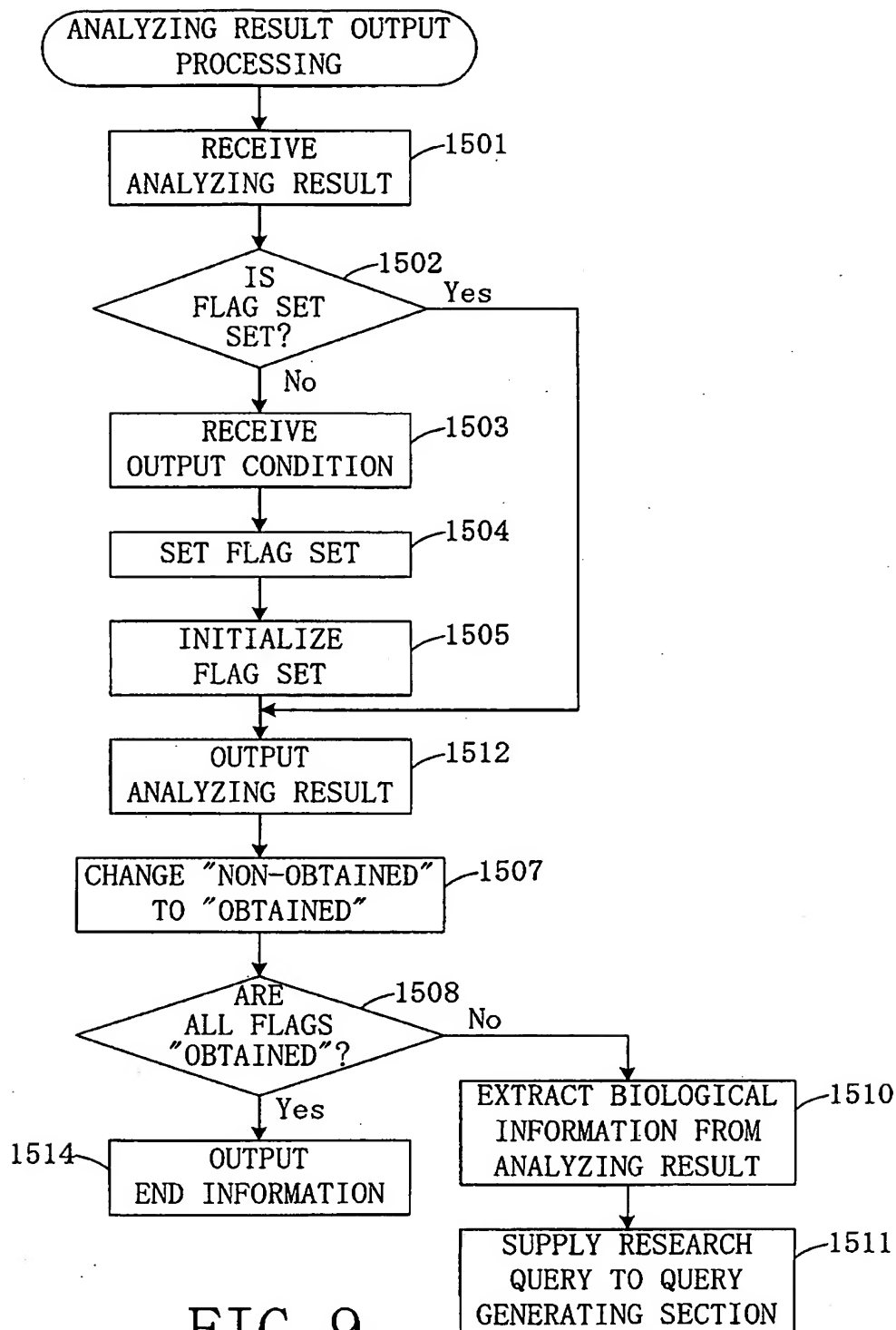


FIG. 9

10/31

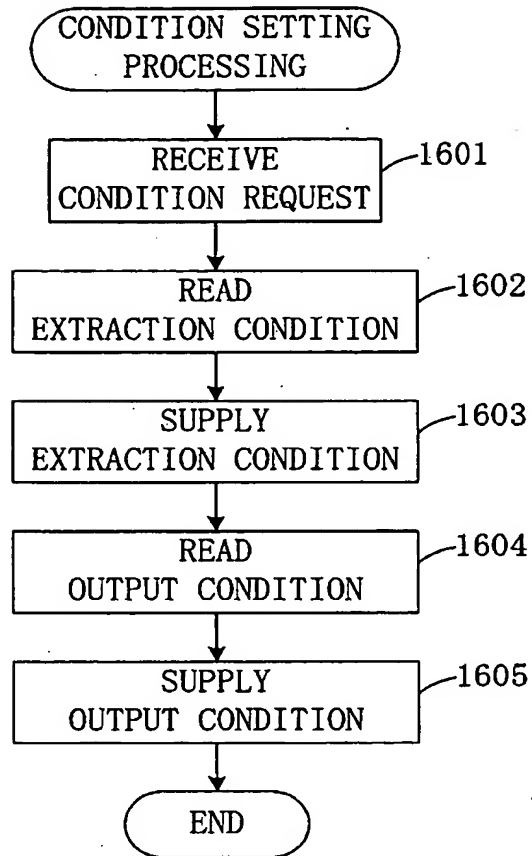


FIG. 10

11/31

Input form

Protein or gene name

Accession No. ☐ protein ☐ gene

Sequence

☐ amino acid ☐ gene

FIG. 11

12/31

EXTRACT AS BYNAME

<input type="checkbox"/> 5172	Hs	SLC26A4; solute carrier family 26, member 4	7q31	■■■■■
	Aliases:	PDS, DFNB4, PENDRIN		
	OMIM:	605648		
	RefSeq[R]:	NM_000441		
	Nucleotide:	AC002467, AF030880		
	Protein:	AAB88773, AAC51873, NP_000432		
<input type="checkbox"/> 23985	Mm	Slc26a4 solute carrier family 26, member 4	12 B1	■ ■■■
	Aliases:	Pds		
	RefSeq:	NM_011867		
	Nucleotide:	AF167411		
	Protein:	AAD51617, NP_035997		
<input type="checkbox"/> 29400	Rn	Slc26a4 solute carrier family 26, member 4	6q16	■ ■■■
	Aliases:	Pds		
	RefSeq:	NM_019214		
	Nucleotide:	AF167412		
	Protein:	AAD51618, NP_062087		
<input type="checkbox"/> 65010	Hs	SLC26A6 solute carrier family 26, member 6	3p21.3	■ ■■■
	Aliases:	DKFZp586E1422		
	RefSeq[R]:	NM_022911, NM_134263, NM_134426		
	Nucleotide:	AB033288, AB102713, AF279265, AF288410, AF416721, AK066237, AL050170, BC017697,		
	Protein:	AAF81911, AAH17697, AAK19153, AAN07094, BAB69041, BAB71126, BAC56861, CAB43308, NP_075062, NP_599025, NP_602298		

FIG. 12

13/31

- | | ACCESSION NUMBER | GI NUMBER |
|------------------------------|--|-----------|
| <input type="checkbox"/> 1. | NT_039548 | |
| | Mus musculus chromosome 12 genomic contig. strain C57BL/6J | |
| | gi 28520880 ref NT_039548.1 Mm12.39588.30[28520880] | |
| <input type="checkbox"/> 2. | NW_043944 | |
| | Rattus norvegicus chromosome 6 WGS supercontig | |
| | gi 26007775 ref NW_043944.1 Rn6.1303[26007775] | |
| <input type="checkbox"/> 3. | NM_134428 | |
| | Homo sapiens solute carrier family 28, member 6 (SLC28A6), transcript variant 3, mRNA | |
| | gi 20338278 ref NM_134428.1[20338278] | |
| | . | |
| | . | |
| | . | |
| <input type="checkbox"/> 35. | A1916698 | |
| | tu89g11 x1 NCLCGAP_Gas4 Homo sapiens cDNA clone IMAGE2258276 3' similar to TRO43511 O43511 PENDRIN. " | |
| | MRNA sequence | |
| | gi 5636553 gb A1916698.1[5636553] | |
| <input type="checkbox"/> 36. | A1747481 | |
| | u115h05. x1 Sugano mouse embryo mewa Mus musculus cDNA clone IMAGE2078921 3' similar to TRO43511 O43511 PENDRIN. " MRNA sequence | |
| | gi 5125725 gb A1747481.1[5125725] | |
| <input type="checkbox"/> 37. | AF030880 | |
| | Homo sapiens pendrin (PDS) mRNA, complete cds | |
| | gi 2654004 gb AF030880.1 AF030880[2654004] | |

FIG. 13

14/31

LOCUS AF030880 4930 bp mRNA linear PR1 01-DEC-1997
DEFINITION Homo sapiens pendrin (PDS) mRNA, complete cds.
ACCESSION AF030880
VERSION AF030880.1 GI:2654004
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 4930)
AUTHORS Ever L.A., Glase, B., Beck, C.J., Idos, J.R., Heym, M.
TITLE Pendred syndrome by mutations in a putative suophate transporter gene (PDS)
JOURNAL Gene 21 (3), 512-532 (1999)
MEDLINE 9750948
PUBMED 893590
REFERENCE 2 (bases 1 to 4920)
AUTHORS Ever L.A., Glase, B., Beck, C.J., Idos, J.R., Heym, M.
TITLE Direct Submission
JOURNAL Submitted (21-SEP-1998) AA Technology Branch, National Institutes

FEATYRES
source

Location/Qualifiers

1..4930

/organism="Homo sapiens"

/db_xref="taxon:9606"

/chromosome="7"

/map="7q22-q31.1"

gene

1..4930

/gene="PDS"

CDS

225..2567

/gene="PDS"

/function="putatibe sulfate transporter"

/note="mutated in Pendred syndrome"

/codon_start=1

/product="pendrin"

/protein_id="AAL57073.1"

/db_xref="GI:2654005"

/translation="MAAPGGRSEPPQLPEYSCSYMVSFVYSELAFQQQHERRLQERK

TLRESLAKCCSCRKRAFGLKTLVPILEWLPKYRVKEWLLSDVISGVSTGLVATLQG

...

EELDIPTKEIEIQVDWNSLPVKVNPVKVPIHSLVLDCCGAISFLDVVGVRSLRVIVKE

FQRIDVNVYFASLQDYVIEKLEQCGFFDDNIRKDTFFLTVHOAILYLQNQVKSQECQG

SILETITLIQDCKDTLELIETELTEEELDVQDEAMRTLAS"

EXTRACT AS BYNAME

EXTRACT AS AMINO
ACID SEQUENCE

BASE COUNT 1454 A 937 C 1082 G 1457 T

ORIGIN

1 ctcagccttc cgggttcggg aaaggggaag aatgcaggag gggtaggatt tctttctga
61 taggatcggg tgggaaagac cgcagcctgt gtgtgtcttt cccttegacc aagggtgtctg
121 ttgtccgta aataaaacgt cccactgcct tctgagagcg ctataaaggc agcggaaggg
.
.
4801 tccagtattg tatatgagtt ttaacaaatt aaaaaatcaa atcatgtaca ttgaaaaata
4861 ttgacacaca tttaaaaaata aatgtaaagt tgtcttttaa actactcgga tgtgtccttt
4921 ctgaacaaaa

EXTRACT AS BASE SEQUENCE

FIG. 14

15/31

BLASTP 2.2.3 [Apr-12-2000]

Sequences producing significant alignments:

Score E
 (bits) Value

ref|NP_000432.1| pendrin [Homo sapiens] >gi|11421915|ref|XP_0049... 1229 0.0
 ref|NP_062087.1| pendrin: Pandred syndrome homolog (human) [Ratt... 1101 0.0
 ref|NP_035997.1| pendrin: Pandred's syndrome [mus musculus] > gi| ... 1098 0.0

...

pir|B82127 sulfate permease family protein VC2031 [imported]... 35 2.8
 emb|CAB62519.1| (AL050358) hypothetical protein [Homo sapiens] 34 3.6
 pir|T16622 hypothetical prtein K02E10.2 - Caenorhabditis alega... 34 4.7

>ref|NP_000432.1| pendrin [Homo sapiens] EXTRACT AS NAME
 ref|XP_004953.1| pendrin [Homo sapiens]
 sp|043511|PEND_HUMAN PENDRIN (SODIUM-INDEPENDENT CHLORIDE/100IDE TRANSPORTER)
 gb|AA51873.1| (AF030880) pendrin [Homo sapiens]
 Length = 780

Score = 1229 bits (3180), Expect = 0.0
 Identities = 649/780 (83%), Positives = 649/780 (83%)

Query: 1 MAAPGGRSEPPQLPEYSCSYMVRPVYSELAFQQQMERRLQERKTLRESLAKCCSCSRKR 60
 MAAPGGRSEPPQLPEYSCSYMVRPVYSELAFQQQHERRLQERKTLRESLAKCCSCSRKR
 Sbjct: 1 MAAPGGRSEPPQLPEYSCSYMVRPVYSELAFQQQHERRLQERKTLRESLAKCCSCSRKR 60

...

Query: 721 TVHDAILYLQNQVKSQEGGGSILETITLIQDCKDXXXXXXXXXXXXXDVQDEAMRTLAS 780
 TVHDAILYLQNQVKSQEGGGSILETITLIQDCKD DVQDEAMRTLAS
 Sbjct: 721 TVHDAILYLQNQVKSQEGGGSILETITLIQDCKDTLELIELTETEEELDVQDVEAMRTLAS 780

FIG. 15

16/31

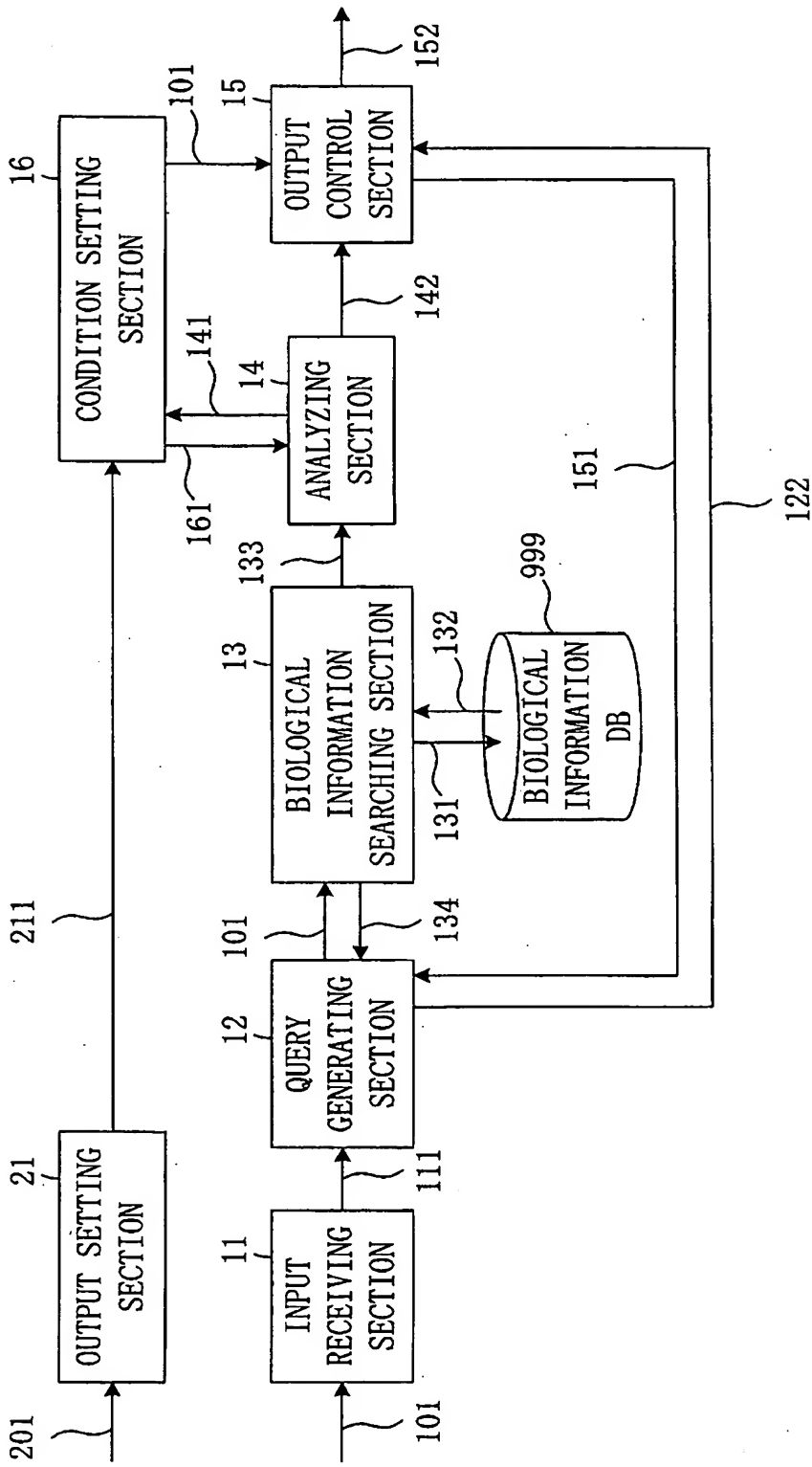


FIG. 16

17/31

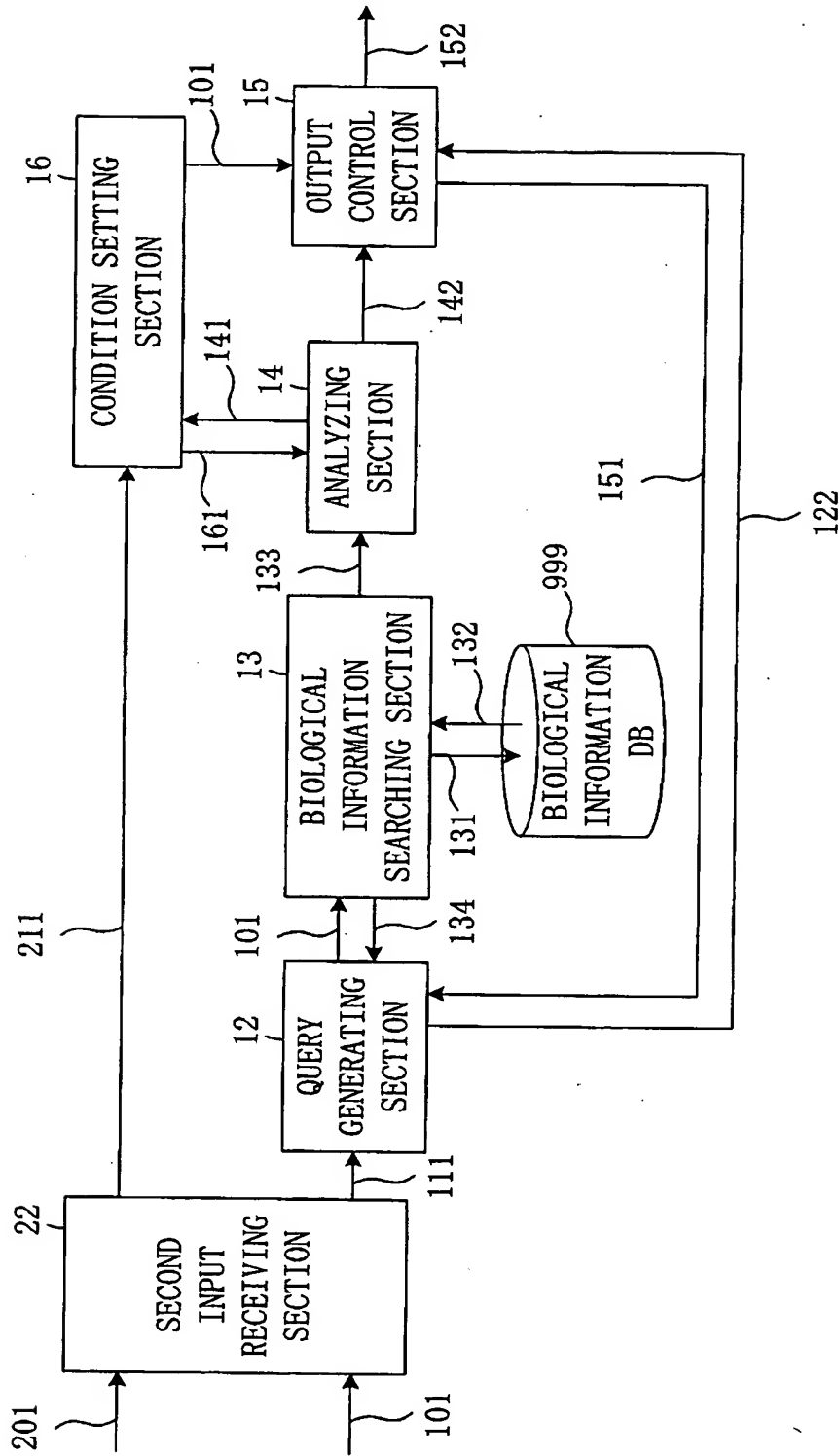


FIG. 17

Entrez ☒ name ☐ Accession No. ☐ Sequence

LocusLink ☒ name ☐ Accession No. (RefSeq) ☐ Accession No. (OMIM) ☐ Accession No. (Entrez) ☐ chromosome position

BLAST ☒ name ☐ Accession No. ☐ Sequence — e-value

submit reset

FIG. 18

19/31

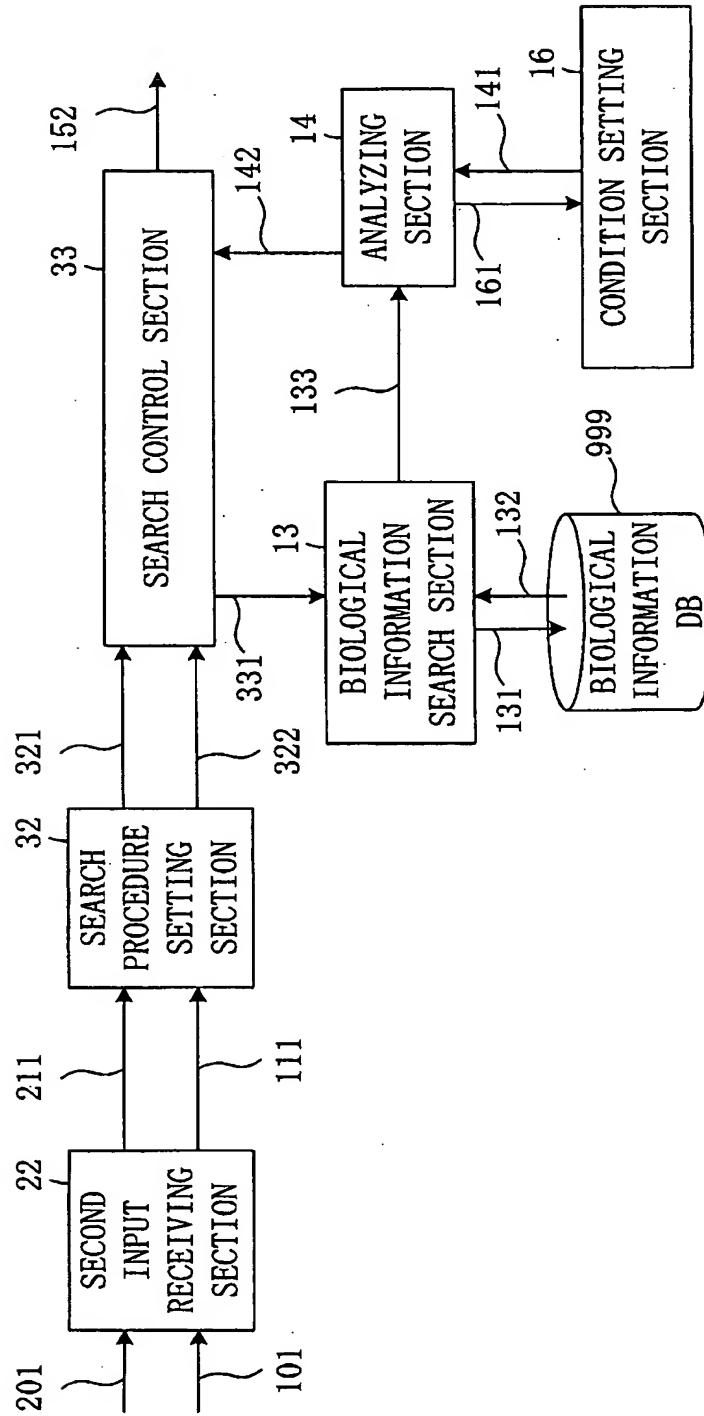


FIG. 19

20/31

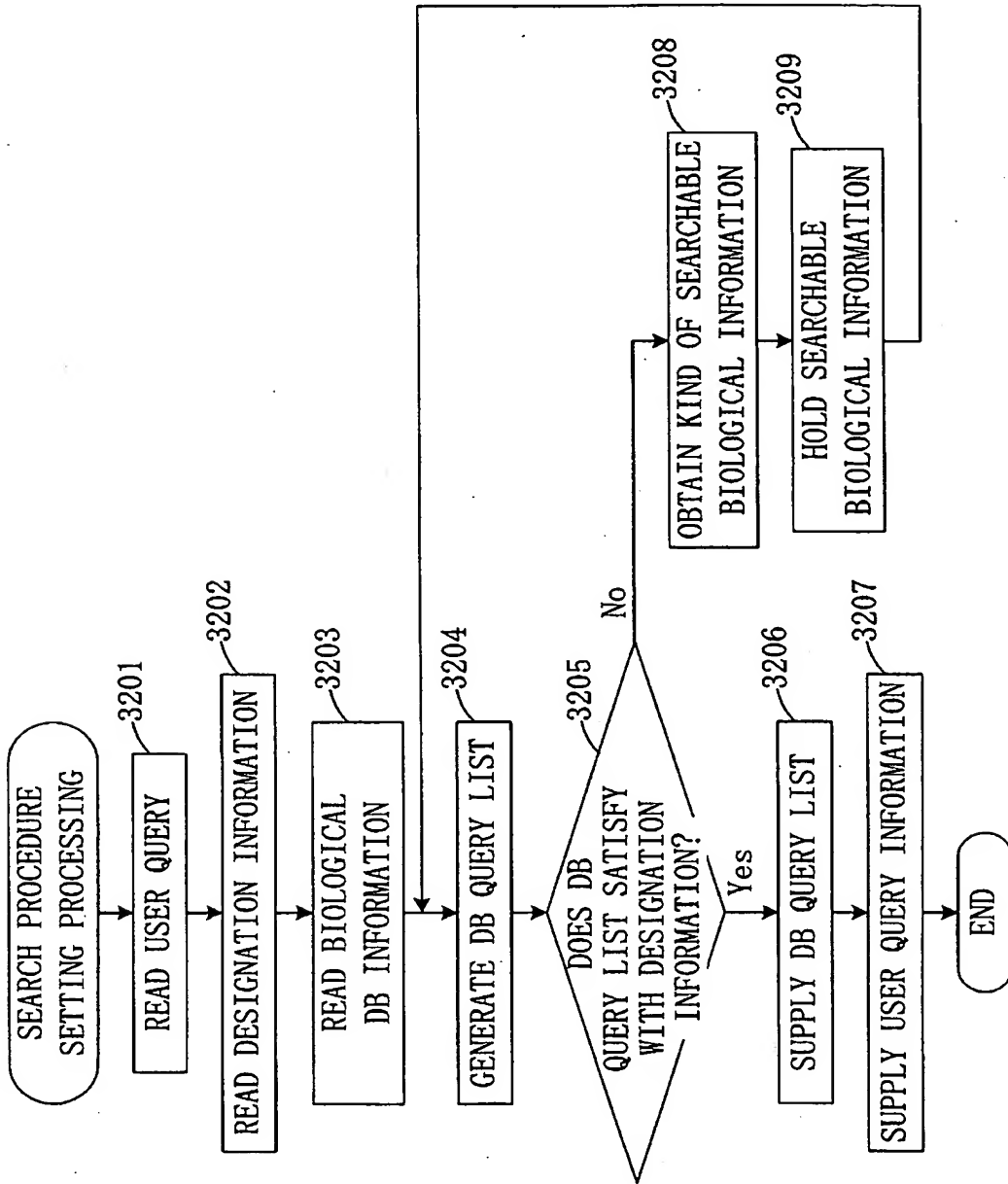


FIG. 20

21/31

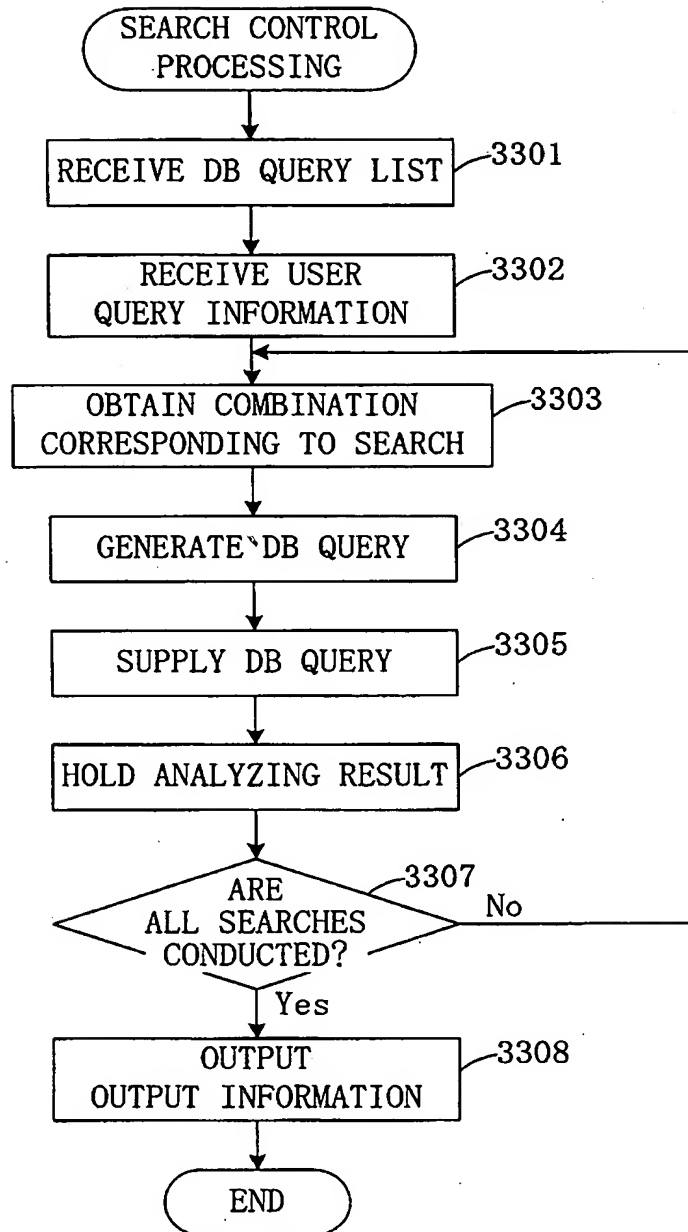


FIG. 21

22/31

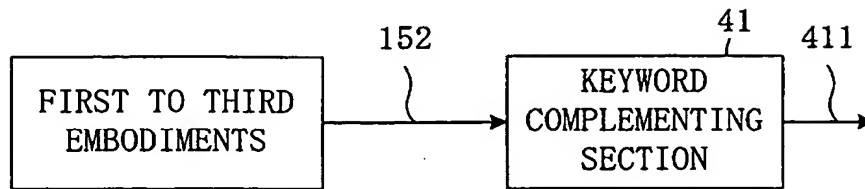


FIG. 22

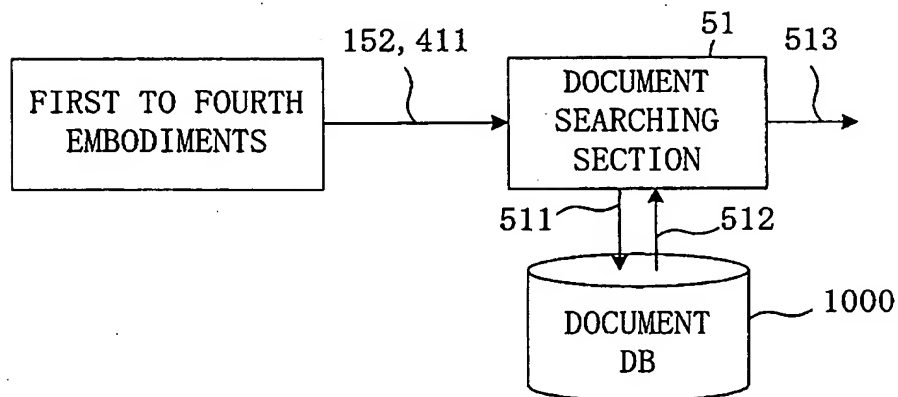


FIG. 23

23/31

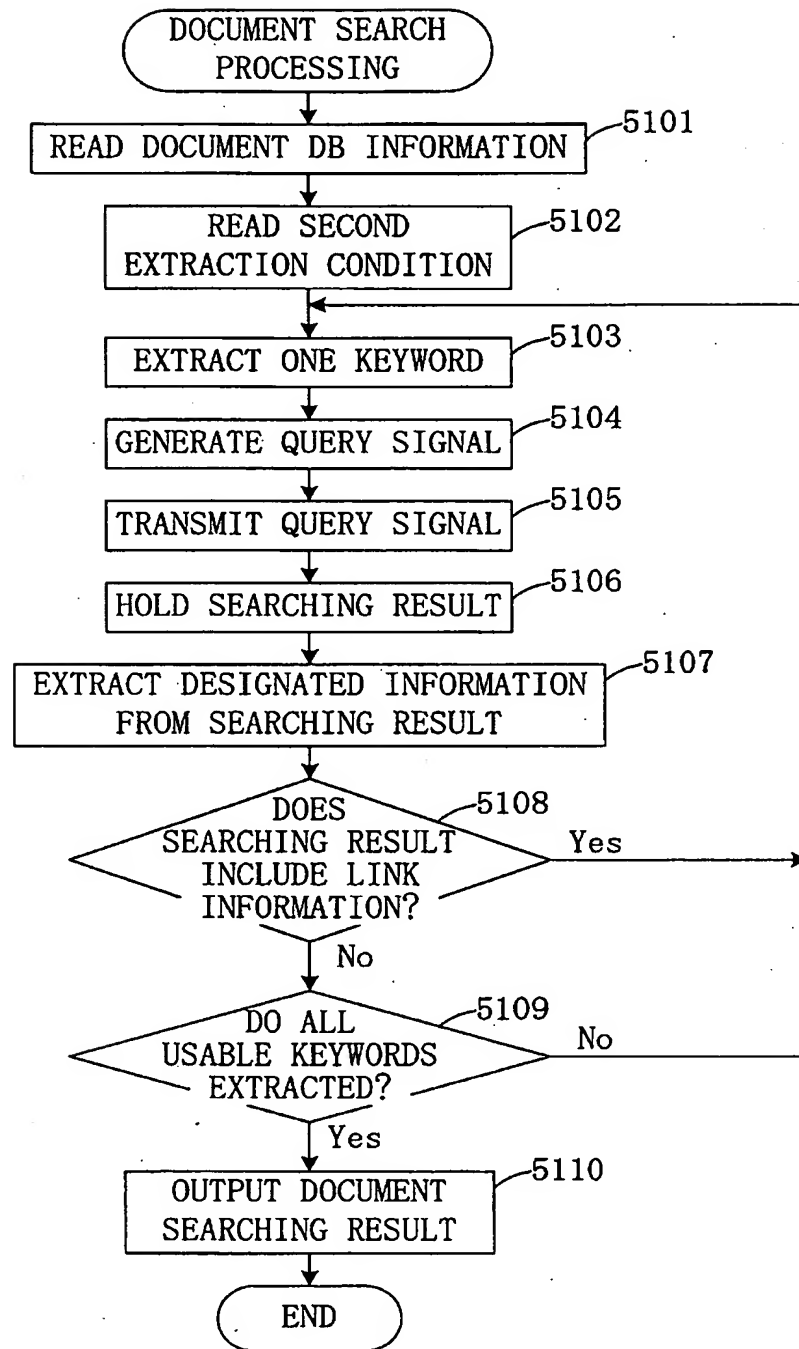


FIG. 24

24/31

ICON	AUTHOR	TITLE	JOURNAL NAME, ISSUED DATE	ID NUMBER
<input type="checkbox"/>	1: Huland M, Kiern AG, Blomg RS, Coal P, Samuel JJ, Johan BR, Steel PK, Enerb K	Lack of pendrin expression leads to ... Development. 2000 Jun; 1(30 (6): 2013-2020 PMID: 12745323 [PubMed - as supplied by publisher]		
<input type="checkbox"/>	2: Grata B, Bieber J, Kuida S	Neonatal thyroid disorders. Horm Res. 2002; 83 Suppl 1: 32-8 PMID: 13284723 [PubMed - in process]		
<input type="checkbox"/>	3: Friesche M, Kwan HT, Froki M, Mad MK, Mielson A	Regulated expression of pendrin in rat. ... Physiol J. 2002 Apr; 283 (4): F384-83 PMID: 12444733 [PubMed - indexed for MEDLINE]		
<input type="checkbox"/>	4: Kono T, Nakagawa N, Suzuki M, Murakami S, Minami A, Kawai A, Kazawa R	Expression of human pendrin in ... J Histochem Cytochem. 2000 Jun; 48 (4): 137-42 PMID: 12487437 [PubMed - indexed for MEDLINE]		
<input type="checkbox"/>	5: Zama G, Bruna L, Knau H, Wang K, Fujii M, Ginzl D, Duh YH, Clark PD	Increasing the effectiveness of radioactive iodine therapy in ... Surgery. 2003 Dec; 133 (5): 885-94; discussion 890. PMID: 12473333 [PubMed - indexed for MEDLINE]		
.				
.				
.				

FIG. 25

25/31

<input type="checkbox"/> 1. Development. 2000 Jun. 1;130 (9). 2013-2020	JOURNAL NAME
Lack of pendrin expression leads to	TITLE
Huland M, Kiern AG, Blomq RS, Coal P, Samuel JJ, Johan BR, Steel PK, Enerb K	AUTHOR
Medical Genetics, Department of Biology, Institute of Biochemistry, Gomsorg University, Box 320, SE-240 30 Gomsorg, Sweden. MRC Institute of Research University Park, York, YK2 4RD, UK. Institute of Anatomy and Cell Biology. —	
Mice that lack the winged helix/forhead gene —	SUMMARY
PMID:12748323 [PubMed - as supplied by publisher]	

FIG. 26

26/31

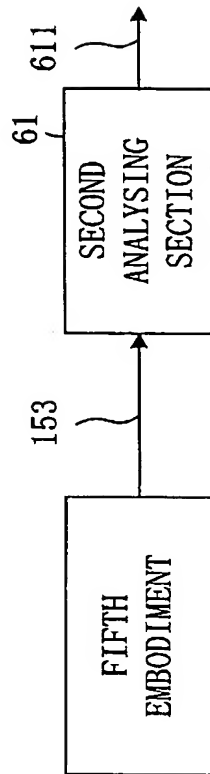


FIG. 27

27/31

<p><input type="checkbox"/> 1. Huland M, Kiem AG, Blom RS, Coal P, Samuel JJ, Johan BR, Steel PK, Enerb K. Lack of pendrin expression leads to ... Development. 2000 Jun; 1130 (6): 2013-2020 PMID: 12748323 [PubMed - as supplied by publisher]</p>	<p>EXAMPLE OF SCHOLARLY PAPER OBTAINED BY ONLY SEARCHING RESULT DUE TO PENDRIN</p>
<p><input type="checkbox"/> 2. Grote B, Bieber J, Knude S. Neonatal thyroid disorders. Horm Res. 2002; 83 Suppl 1: 32-8 PMID: 22284723 [PubMed - in process]</p>	<p>EXAMPLE OF SCHOLARLY PAPER OBTAINED BY BOTH SEARCHING RESULT DUE TO PENDRIN AND SEARCHING RESULT DUE TO SLC26A4</p>
<p><input type="checkbox"/> 12: Billerme NS, Hill AJ. Prolactin regulation of the pendrin⁻ iodide transporter in ... Physiol J. 2003 Feb; 322 (2): E34-8. PMID: 12384373 [PubMed - indexed for MEDLINE]</p>	
<p><input type="checkbox"/> 13: Kamiński JA, Wang G, Enerb J, Green PE, Giebs G, Axon SO. Formate-stimulated NaCl absorption in ... of the pendrin protein. Physiol J. 2003 Nov; 331 (4): F874-83. PMID: 12248297 [PubMed - indexed for MEDLINE]</p>	
<p><input type="checkbox"/> 14: Rot PP, Hirscheberg M, Manu S, Sasaki K, Roy RI, Green DE, Kon K, Lipin J, Yen MP. Retention of pendrin in ... Hum Genet. 2002 Jan; 121 (45): 2813-32. PMID: 12157323 [PubMed - indexed for MEDLINE]</p>	

FIG. 28

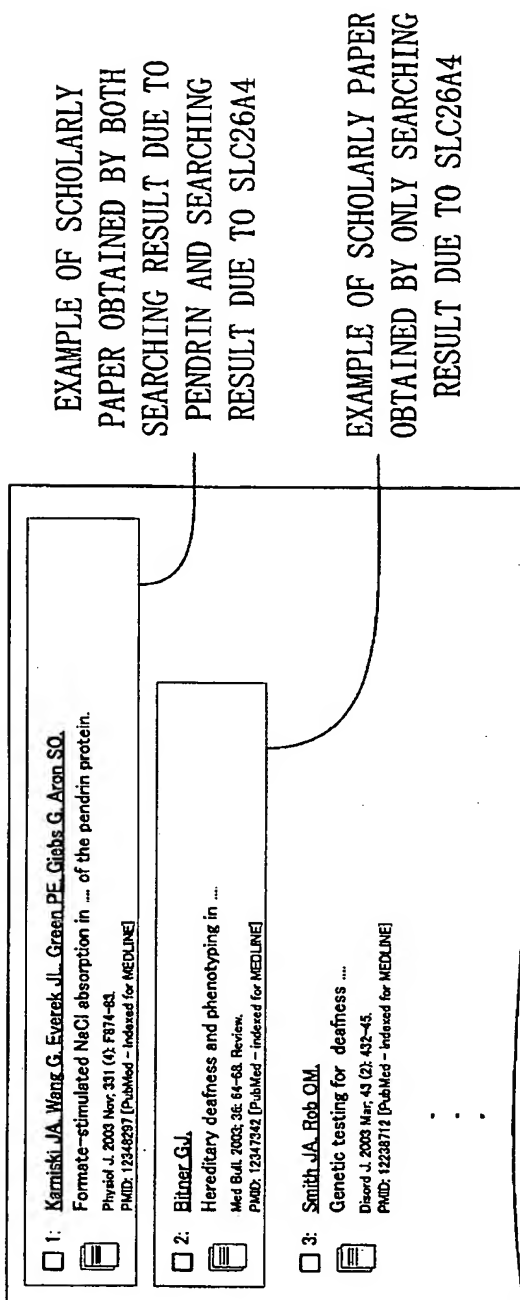


FIG. 29

29/31

EXAMPLE OF SCHOLARLY PAPER
OBTAINED BY ONLY SEARCHING
RESULT DUE TO PENDRIN

EXAMPLE OF SCHOLARLY
PAPER OBTAINED BY BOTH
SEARCHING RESULT DUE TO
PENDRIN AND SEARCHING
RESULT DUE TO SLC26A4

EXAMPLE OF SCHOLARLY PAPER
OBTAINED BY ONLY SEARCHING
RESULT DUE TO SLC26A4

☐ 1. Huland M, Kiern AG, Blomn RS, Coal P, Samuel JJ, Johan BR, Steel PK, Enerb K.
Lack of pendrin expression leads to
Development. 2000 Jun; 1:130 (9): 2013-2020
PMID: 12748323 [PubMed - as supplied by publisher]

☐ 2: Grata B, Bleher J, Kaude S.
Neonatal thyroid disorders.
Horm Res. 2002; 83 Suppl 1: 32-8
PMID: 23284733 [PubMed - in process]

☐ 12: Billime NS, Hill AJ.
Prolactin regulation of the pendrin- iodide transporter in ...
Physiol J. 2003 Feb; 322 (2): E34-8.
PMID: 12384373 [PubMed - indexed for MEDLINE]

☐ 13: Kaminski JA, Wang G, Everek JL, Green PE, Giebs G, Anon SO.
Formate-stimulated NaCl absorption in ... of the pendrin protein.
Physiol J. 2003 Nov; 331 (4): F874-83.
PMID: 12348287 [PubMed - indexed for MEDLINE]

☐ 14: Rot PP, Hirscheberg M, Maru S, Sasaki K, Roy RL, Green DE, Kon K, Lippin J, Yen MP.
Retention of pendrin in ...
Hum Genet. 2002 Jan; 1: 23 (45): 2813-32.
PMID: 12357323 [PubMed - indexed for MEDLINE]

☐ 15: Bitner GJ.
Hereditary deafness and phenotyping in ...
Med Bull. 2003; 36: 64-68. Review.
PMID: 12347342 [PubMed - indexed for MEDLINE]

FIG. 30

30/31

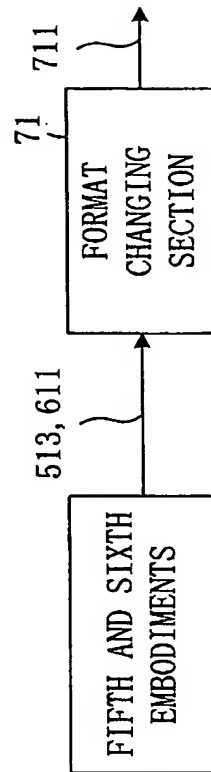


FIG. 31

31/31

<p>***KEYWORD*** pendrin</p> <p>***Another Name*** [GenBank] •PDS [LocusLink] •SLC26A4 •solute carrier family 26, member 4 •PDS •DFNB4 •SLC26A6 •solute carrier family 26, member 6 •DKFZp586B1442 [BLAST] •down regulated in adenoma •down regulated in adenoma protein •sulfate anion transporter 1 •diastrophic dysplasia</p>	<p>Thyroid 2002 Jan; 12 (3): 849-54</p> <p>Expression of nicotinamide adenine dinucleotide phosphata oxidase</p> <p>Lacro J, Nocer M, Minor G, Cailla J, Viciore B, David C, Fiechi M, Bidart GP. Department of Biochemistry, Institute of Medical Genetics, France.</p> <p>Diox2, and probably Diox1 are ...</p> <p>... of sodium iodide symporter (NIS), pendrin and ...</p> <p>PMID: 11738123 [PubMed - indexed for MEDLINE]</p> <p>Thyroid 2002 Jan; 12 (3): 849-54</p> <p>Update on intrathyroidal iodine</p> <p>Dun A, Samuel EL Department of Medicine, Institute of Biochemistry, USA. dune@xxx.edu</p> <p>The thyroid concentrates iodide from the serum and ...</p>
--	---

DISPLAY SUMMARY
OF DOCUMENT IN
FORM OF LIST

CHANGE DISPLAY
COLOR OF KEYWORD

DISPLAY GENE OR
PROTEIN BYNAME

FIG. 32